Amendments to the Claims:

- 1. to 28. (Cancelled)
- 29. (Currently amended) A method of immunizing a host against infection caused by a strain of *Chlamydia*, which comprises:

administering to the host an immunoeffective amount of an attenuated bacteria harbouring a vector comprising a nucleic acid molecule encoding at least one immunoprotection-inducing *Chlamydia* protein or a fragment thereof which generates a *Chlamydia* protein specific immune response and a promoter operatively coupled to said nucleic acid molecule for expression of said *Chlamydia* protein or fragment thereof in cells of the host but not in said attenuated bacteria.

- 30. (Original) The method of claim 29 wherein said immunoprotection inducing *Chlamydia* protein or fragment thereof is a major outer membrane protein (MOMP) of a strain of *Chlamydia*.
- 31. (Original) The method of claim 30 wherein said strain of *Chlamydia* is a strain of *Chlamydia pneumoniae*.
- 32. (Original) The method of claim 30 wherein said strain of *Chlamydia* is a strain of *Chlamydia trachomatis*.
- 33. (Cancelled)
- 34. (Currently amended) The method of claim <u>29</u> 33 wherein said promoter is a cytomegalovirus promoter.
- 35. (Currently amended) The method of claim 29 33 wherein said vector is a plasmid vector.
- 36. (Currently amended) The method of claim 35 wherein said plasmid vector <u>is</u> has the identifying characteristics of pcDNA3/MOMP as seen in Figure 5.
- 37. (Original) The method of claim 29 wherein said attenuated bacteria is an attenuated strain of Salmonella.

- 38. (Original) The method of claim 37 wherein said attenuated strain of *Salmonella* is an attenuated strain of *Salmonella typhimurium*.
- 39. (Original) The method of claim 29 wherein said administration is effected to mucosal surfaces.
- 40. (Original) The method of claim 39 wherein said administration is effected by intranasal administration.